Rare Disease Medications

	Membe	r and Medicat	ion Informatio	n (required)	
Me	ember ID:		Member Name:		
DOB:			Weight:		
Medication Name/ Strength:			Dose:		
Di di di					
ווט	rections for use:				
Provider Information (required)					
Name:		NPI:		Specialty:	
Contact Person:		Office Phone:		Office Fax:	
	FAX FORM AND RELEVA				
	CHART NOTES and/or U				
	eria for Approval (all criteria must be m				
	Medication is prescribed by or in consul • Specialist name and credentials		· ·		
	Documented diagnosis: Chart note page #:				
	Genetic testing, if applicable. Chart note page #:				
Other confirmation testing, if applicable. Chart note page #:					
	 Use must follow FDA-approved labeling (including monitoring for boxed warnings and contraindications). Applicable monitoring for boxed warnings. Chart note page #:				
	If current treatment standards recomme			ons prior to use of the requested drug,	
	document the use of appropriate first line. Treatment/Interventions:			page #:	
_	Off Label or Compendia Use Additional Criteria: Requests for any off-label indications must be supported by at least one (1) major multi-site study or three (3) smaller studies published in JAMA, NEJM, Lancet or other peer review specialty medical				
	journals within the most recent five (5) years. Supporting documentation must be included. Compendia use must be recommended by generally-accepted compendia such as American Hospital Formulary Service Drug Information (AHFS), United				
	States Pharmacopeia-Drug Information (USP-DI), the DRUGDEX Information System, and the peer-reviewed medical literature.				
	Additional drug-specific criteria may apply and may guide adjudication; additional information may be needed. See addendu				
	for details, page 2.				
	authorization Criteria, if applicable: lated letter of medical necessity or upda	ted chart notes demor	nstrating positive clinica	al response	
	ial Authorization: Up to six (6) months, it uthorization: Up to one (1) year, if applie				
Not	e:				
	Use appropriate HCPCS code for bill Coverage and Reimbursement and		rb utah gay/stalan/laal	uun/Coveragel eeluun nha	
	Coverage and Reimbursement code HCPCS NDC Crosswalk: https://heal	•			
PRC	OVIDER CERTIFICATION				
I he	reby certify this treatment is indicated, n	necessary and meets th	ne guidelines for use.		
Pres	scriber's Signature		Date		

UTAH DEPARTMENT OF HEALTH, PRIOR AUTHORIZATION REQUEST FORM

Brand (generic)	Additional Criteria	
Luxturna (voretigene)	What is the patient's diagnosis:	
	☐ Biallelic RPE65 mutation-associated retinal dystrophy	
	Other, please specify:	
	Which eye is being treated:	
	☐ Left eye ☐ Right eye ☐ both eyes	
	*If BOTH eyes, do you agree that the initial eye's injection and the second eye's injection will be	
	administered at least 6 days apart?	
	☐ Yes ☐ No	
	Has the patient received Luxturna previously?	
	Yes No	
	*If yes, which eye(s) were previously treated?	
	☐ Left eye ☐ Right eye	
	*If treating the additional eye, do you agree that the initial eye's injection and the second eye's	
	injection will be administered at least 6 days apart? ☐ Yes ☐ No	
	Tes Till	
	Documented diagnosis of biallelic RPE65 mutation-associated retinal dystrophy confirmed by	
	genetic testing? (please include genetic testing results)	
	Yes No	
	Authorization: once per lifetime	
Zolgensma	Authorization: once per lifetime	